Annual Index

Compiled by Roseann Caffaro

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Edited by Roseann Caffaro

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Mandelamine® (methenamine mandelate)

CAUTION: Federal law prohibits dispensing without prescription.

Description. Mandelamine, a urinary antibacterial agent, is the chemical combination of mandelic acid with methenamine.

Indications. Mandelamine (methenamine mandelate) is indicated for the suppression or elimination of bacteriuria associated with pyelonephritis, cystitis, and other chronic urinary tract infections; also for infected residual urine sometimes accompanying neurologic diseases. When used as recommended, Mandelamine is particularly suitable for long-term therapy because of its safety and because resistance to the nonspecific bactericidal action of formaldehyde does not develop. Pathogens resistant to other antibacterial agents may respond to Mandelamine because of the nonspecific effect of formaldehyde formed in an acid urine.

Contraindications. Contraindicated in renal insufficiency.

Precautions. Dysuria may occur (usually at higher than recommended dosage). This can be controlled by reducing the dosage and the acidification. When urine acidification is contraindicated or unattainable (as with some urea-splitting bacteria), the drug is not recommended.

To avoid inducing lipid pneumonia, administer Mandelamine Suspension Forte and Mandelamine Suspension with care to elderly, debilitated or otherwise susceptible patients.

Adverse Reactions. An occasional patient may experience gastrointestinal disturbance or a generalized skin rash.

Dosage and Management. The average adult dosage is 4 grams daily given as 1.0 gram after each meal and at bedtime. Children 6 to 12 should receive half the adult dose and children under 6 years of age should receive 250 mg per 30 lb body weight, four times daily. (See chart.) Since an acid urine is essential for antibacterial activity with maximum efficacy occurring at pH 5.5 or below, restriction of alkalinizing foods and medication is thus desirable. If testing of urine pH reveals the need, supplemental acidification should be given.

Mandelamine Dosages	ADULTS	CHILDREN
Tablets and Granules		
1.0 gram	1 tablet q.i.d.	—
	1 packet* q.i.d.	—
0.5 gram	2 tablets q.i.d.	(Ages 6-12) 1 tablet q.i.d.
		1 packet* q.i.d.
0.25 gram	—	(Age under 6) 1 tablet per 30 lb body weight q.i.d.
Suspension Forte		
500 mg/5 ml teaspoonful	2 teaspoonfuls (10 ml) q.i.d.	(Ages 6-12) 1 teaspoonful (5 ml) q.i.d.
Suspension		
250 mg/5 ml teaspoonful	—	(Age under 6) 1 teaspoonful (5 ml) per 30 lb body weight q.i.d.

*Contents of packet to be dissolved in 2-4 oz water immediately before using.
Shake Suspensions well before using.

STORE BETWEEN 59° and 86°F (15° and 30°C).

Supplied: 1.0 gm Tablets W/C 172: purple, enteric coated in bottles of 100 (N 0047-0172-51) and 1000 (N 0047-0172-60); also unit dose in 10 x 10 strips (N 0047-0172-11).

Granules (1.0 gm): orange-flavored individual packets; cartons of 56 (N 0047-0176-11).

0.5 gm Tablets W/C 171: brown, enteric coated in bottles of 100 (N 0047-0171-51) and 1000 (N 0047-0171-60); unit dose in 10 x 10 strips (N 0047-0171-11).

Granules (0.5 gm): orange-flavored individual packets; cartons of 56 (N 0047-0177-11).

0.25 gm Tablets W/C 170: brown, enteric coated in bottles of 100 (N 0047-0170-51) and 1000 (N 0047-0170-60).

Suspension Forte,† 500 mg/5 ml teaspoonful: pink, cherry-flavored in bottles of 8 fl oz (N 0047-0174-08) and 16 fl oz (N 0047-0174-16). Unit Dose – 10 ml (N 0047-0174-10). U.S. Patent No. 3,077,438.

Suspension,† 250 mg/5 ml teaspoonful: cream-colored, coconut-flavored in bottles of 4 fl oz (N 0047-0173-04) and 16 fl oz (N 0047-0173-16).

Full information is available on request. M-GP-61-4/c

† Suspensions are in vegetable oil. Shake well before using.

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